**PURPLE HAND SANITIZER- benzalkonium chloride 0.13% spray**
**ASN Technologies Inc**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**Purple Hand Sanitizer**

## Active Ingredient

Benzalkonium Chloride 0.13%

## Purpose

Antiseptic

## Uses

To help reduce bacteria that potentially can cause disease. For when soap and water are not available.

## Warnings

For external use only.

## Do Not Use

on open skin wounds or on children less than two months of age

## When using this product

keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

## Stop use and ask a doctor

if irritation or rash occurs. These may be signs of a serious condition.

## Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

## Directions

On dry hands, place enough product in your palm to thoroughly cover your hands. Rub hands together briskly until hands are entirely dry. Supervise children under 6 years of age when using this product to avoid swallowing.

## Other Information

Store between 15-30C (59-86F)
Avoid heating and temperatures above 40C (104F)

## Inactive Ingredients

Silica complex 1-octadecanaminium, N,n-dimethyl-n-(3-(trihydroxysilyl)propyl) Chloride, Polyoxyethylenesorbitan monolaurate, Glycerol, Sodium Benzoate, Aqua

## Purple Hand Sanitizer


| PURPLE HAND SANITIZER | |
|---|---|
| benzalkonium chloride 0.13% spray | |

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:82294-001 |
| **Route of Administration** | TOPICAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **BENZALKONIUM CHLORIDE** (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y) | BENZALKONIUM CHLORIDE | 0.078 g in 60 mL |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **SODIUM BENZOATE** (UNII: OJ245FE5EU) | |
| **POLYSORBATE 20** (UNII: 7T1F30V5YH) | |
| **OCTADECYLDIMETHYL(3-TRIHYDROXYSILYLPROPYL)AMMONIUM CHLORIDE** (UNII: GLJ50K866T) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:82294-001-60 | 60 mL in 1 BOTTLE; Type 0: Not a Combination Product | 12/01/2021 | |
| **2** | NDC:82294-001-40 | 40 mL in 1 BOTTLE; Type 0: Not a Combination Product | 12/01/2021 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part333A | 12/01/2021 | |

## Labeler - ASN Technologies Inc (204344589)

## Registrant - ASN Technologies Inc (204344589)

Revised: 12/2021 ASN Technologies Inc